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CLAIMS:

1. A process for synthesising hydrocarbons, which process includes

feeding a gaseous feedstock comprising hydrogen, carbon monoxide and carbon dioxide, into a dimethyl ether (DME) synthesis stage, the gaseous feedstock having a syngas number (SN) between 1.8 and 2.2, where

SN =
$$[H_2] - [CO_2]$$

10 - $[CO] + [CO_2]$

and where [H₂], [CO] and [CO₂] respectively are the molar proportions of hydrogen, carbon monoxide and carbon dioxide in the gaseous feedstock;

in the DME synthesis stage, converting a portion of the gaseous feedstock into a DME product and gaseous products;

separating the DME product from unreacted gaseous reactants and the gaseous products to obtain a tail gas comprising hydrogen and carbon monoxide and carbon dioxide;

recycling a portion of the tall gas from the DME synthesis stage to the DME synthesis stage, a ratio of tall gas recycle to gaseous feedstock being between about 0: 1 and about 2:1;

feeding the tail gas Into a Fischer-Tropsch hydrocarbon synthesis stage, which is a two-phase high temperature catalytic Fischer-Tropsch hydrocarbon synthesis stage; and

allowing the hydrogen, carbon monoxide and carbon dioxide at least partially to react catalytically in the Fischer-Tropsch hydrocarbon synthesis stage to form hydrocarbons, the hydrocarbons formed in the Fischer-Tropsch hydrocarbon synthesis stage thus being gaseous hydrocarbons at the operating pressure and temperature of the Fischer-Tropsch hydrocarbon synthesis stage.

2. The process as claimed in claim 1, in which converting a portion of the gaseous feedstock into a DME product and gaseous products includes contacting the gaseous feedstock with a catalyst or catalysts that enhance methanol synthesis and methanol dehydration reactions.

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3. The process as claimed in claim 1 or claim 2, in which the DME product includes a mixture of DME and methanol and which includes converting the DME product into light olefins in a light olefins production stage without increasing the DME concentration in the DME product.

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4. The process as claimed in any one of the preceding claims, in which the DME synthesis stage is operated at a pressure of between about 50 bar(g) and about 100 bar(g) to ensure that overall CO + CO₂ conversion in the DME synthesis stage is between about 20 % and about 80 %.

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5. The process as claimed in any one of the preceding claims, which includes recycling some of the Fischer-Tropsch hydrocarbon synthesis stage tail gas to the Fischer-Tropsch hydrocarbon synthesis stage, to obtain high overall $CO + CO_2$ conversions in the Fischer-Tropsch hydrocarbon synthesis stage of at least 80 %.

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6. The process as claimed in any one of the preceding claims, which includes recycling some of the Fischer-Tropsch hydrocarbon synthesis stage tail gas to the Fischer-Tropsch hydrocarbon synthesis stage, a ratio of Fischer-Tropsch tail gas recycle to the tail gas from the DME synthesis stage fed to the Fischer-Tropsch hydrocarbon synthesis stage being between 2.5:1 and 1:1.5.

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7. The process as claimed in claim 3, which includes, in a separation stage, separating light hydrocarbons from the Fischer-Tropsch hydrocarbon synthesis stage tail gas and converting these light hydrocarbons, together with the DME product, into light olefins with a carbon number from 2 to 4 in the light olefins production stage.

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8. The process as claimed in claim 3 or claim 7, in which gaseous hydrocarbons and any unreacted hydrogen, unreacted carbon monoxide, and CO₂ are withdrawn from the Fischer-Tropsch hydrocarbon synthesis stage, and separated into one or more condensed liquid hydrocarbon streams, a reaction water stream and a Fischer-Tropsch hydrocarbon synthesis stage tail gas, the process further including treating the condensed liquid hydrocarbons from the Fischer-Tropsch hydrocarbon synthesis stage, to provide a light hydrocarbon fraction, including naphtha, which is converted, together

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with the DME product, in the light olefin production stage to light olefins, and to provide a diesel fraction.

- 9. The process as claimed in claim 3 or claim 7 or claim 8, which includes using separation equipment to recover C_2 - C_4 light olefins from the Fischer-Tropsch hydrocarbon synthesis stage and in which C_2 - C_4 light olefins from the light olefins production stage are recovered using the same separation equipment that is used to recover the C_2 - C_4 light olefins produced by Fischer-Tropsch synthesis.
- 10. The process as claimed in any one of the preceding claims, in which the twophase high temperature catalytic Fischer-Tropsch Hydrocarbon synthesis stage employs an iron catalyst.

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